

## WEST Search History





DATE: Tuesday, June 01, 2004

Hide?	Set Name	Query	Hit Count
	<i>DB=PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=YES; OP=ADJ</i>		
<input type="checkbox"/>	L50	L39 and (fluid with rout\$4)	1
<input type="checkbox"/>	L49	L43 and (fluid with rout\$4)	1
<input type="checkbox"/>	L48	L43 and ((dynamic\$4 or electric\$4) with field with gradient with focus\$4)	3
	<i>DB=USPT,PGPB,JPAB,EPAB,DWPI,TDBD; PLUR=YES; OP=ADJ</i>		
<input type="checkbox"/>	L47	6504368	4
<input type="checkbox"/>	L46	L39 and (vargas)	1
<input type="checkbox"/>	L45	L44 and ((fused adj silica) or PEEK)	6
<input type="checkbox"/>	L44	L43 and (fluid\$5 with (rout\$4 or control\$4 or direct\$4 or separat\$4 or divert\$5 or select\$6))	6
<input type="checkbox"/>	L43	L42 and (detect\$6 or reception or receiv\$5 or sens\$5)	7
<input type="checkbox"/>	L42	L41 and (site or location or chamber or tube or rout\$4 or fluid\$6)	7
<input type="checkbox"/>	L41	L39 and (((sample or analyte) with (hold\$6 or held)) or void)	7
<input type="checkbox"/>	L40	L39 and ((fused adj silica) or PEEK)	12
<input type="checkbox"/>	L39	L37 and ((microcoil or micro-coil or (micro adj coil)) with (helical\$3 or spiral\$3 or solenoid\$4))	29
<input type="checkbox"/>	L38	L37	173184
<input type="checkbox"/>	L37	((magnetic adj resonance) or MRI or NMR)	173184
<input type="checkbox"/>	L36	L35 and (microcoil with (helical\$3 or spiral\$3 or solenoid\$4))	2
<input type="checkbox"/>	L35	fetzner	200
<input type="checkbox"/>	L34	L33 and (memory)	1
<input type="checkbox"/>	L33	L32 and (photo\$9)	2
<input type="checkbox"/>	L32	L30 and (pump\$4)	2
<input type="checkbox"/>	L31	L30 and (pump)	1
<input type="checkbox"/>	L30	L29 and (sound or sonic\$8 or acoustic\$6 or IR or UV)	2
<input type="checkbox"/>	L29	L27 and (heat\$4)	2
<input type="checkbox"/>	L28	L27 and (gradient)	1
<input type="checkbox"/>	L27	L26 and ((capillary adj electrophoresis) or CE)	2
<input type="checkbox"/>	L26	L25 and (liquid adj chromatography)	2
<input type="checkbox"/>	L25	L24 and (extraction with chamber)	2
<input type="checkbox"/>	L24	L23 and (microcoil with planar)	2
<input type="checkbox"/>	L23	L22 and (microcoil with (helical, pr spiral, or solenoid\$4))	2

<input type="checkbox"/>	L22	L21 and (optimiz\$9)	2
<input type="checkbox"/>	L21	L20 and (dimension\$6 or spectra or spectrum or spectral)	2
<input type="checkbox"/>	L20	L18 and ((fused adj silica) or PEEK)	2
<input type="checkbox"/>	L19	L18 and ((fused adj silica) or PEEK)	1
<input type="checkbox"/>	L18	L17 and (sample with size)	2
<input type="checkbox"/>	L17	L16 and ((one or "1") with dimension\$6)	4
<input type="checkbox"/>	L16	L14 and (data with processing)	4
<input type="checkbox"/>	L15	L14 and (controller)	2
<input type="checkbox"/>	L14	6194900	10
<input type="checkbox"/>	L13	L11 not L12	5
<input type="checkbox"/>	L12	L11 and (coil or probe)	11
<input type="checkbox"/>	L11	L9 and (fused with silica)	16
<input type="checkbox"/>	L10	L9 and (microcoil or micro-coil or "micro coil")	0
<input type="checkbox"/>	L9	L8 and (PEEK or polyetheretherketone or poly-ether-ether-ketone)	119
<input type="checkbox"/>	L8	L1 and (polytetrafluoroethylene or poly-tetra-fluoro-ethylene)	3044
<input type="checkbox"/>	L7	L3 and (polytetrafluoroethylene or poly-tetra-fluoro-ethylene)	0
<input type="checkbox"/>	L6	L4 and (polytetrafluoroethylene or poly-tetra-fluoro-ethylene)	1
<input type="checkbox"/>	L5	L4 and (polytetrafluoroethylene or poly-tetra-fluoro-ethylene)	0
<input type="checkbox"/>	L4	L3 and (fused with silica)	3
<input type="checkbox"/>	L3	L2 and (microcoil or micro-coil or "micro coil")	6
<input type="checkbox"/>	L2	L1 and (PEEK or polyetheretherketone or poly-ether-ether-ketone)	527
<input type="checkbox"/>	L1	((magnetic adj resonance) or MRI or NMR)	173184

END OF SEARCH HISTORY

## Hit List

Search Results - Record(s) 1 through 3 of 3 returned.

☐ 1. Document ID: US 20020149369 A1

Using default format because multiple data bases are involved.

L4: Entry 1 of 3

File: PGPB

Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020149369

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020149369 A1

TITLE: Microfluidic device with multiple microcoil NMR detectors

PUBLICATION-DATE: October 17, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	
Norcross, Jim	Champaign	IL	US	
Strand, David	Sherborn	MA	US	
Sweedler, Jonathan	Urbana	IL	US	

US-CL-CURRENT: 324/321; 324/306, 435/4

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	NMC	Drawings
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☐ 2. Document ID: US 6061587 A

L4: Entry 2 of 3

File: USPT

May 9, 2000

US-PAT-NO: 6061587

DOCUMENT-IDENTIFIER: US 6061587 A

TITLE: Method and apparatus for use with MR imaging

DATE-ISSUED: May 9, 2000

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Kucharczyk; John	Edina	MN		
Moseley; Michael E.	Redwood City	CA		

US-CL-CURRENT: 600/411, 600/431, 600/432, 600/433, 604/151, 604/152, 604/153,  
604/154, 604/155, 604/21, 604/93.01

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWIC	Drawings
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☐ 3. Document ID: US 6026316 A

L4: Entry 3 of 3

File: USPT

Feb 15, 2000

US-PAT-NO: 6026316

DOCUMENT-IDENTIFIER: US 6026316 A

**\*\* See image for Certificate of Correction \*\***

TITLE: Method and apparatus for use with MR imaging

DATE-ISSUED: February 15, 2000

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Kucharczyk; John	Edina	MN		
Moseley; Michael E.	Redwood City	CA		

US-CL-CURRENT: 600/420; 324/309

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWIC	Drawings
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Clear	Generate Collection	Print	Fwd Refs	Bkwd Refs	Generate OACS
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Term	Documents
FUSED	231920
FUSED S	0
SILICA	436960
SILICA S	16978
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(L3 AND (FUSED WITH SILICA)).USPT,PGPB,JPAB,EPAB,DWPI,TDBD.	3

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[Generate OACS](#)

Search Results - Record(s) 1 through 7 of 7 returned.

☐ 1. Document ID: US 20020149369 A1

Using default format because multiple data bases are involved.

L43: Entry 1 of 7

File: PGPB

Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020149369

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020149369 A1

TITLE: Microfluidic device with multiple microcoil NMR detectors

PUBLICATION-DATE: October 17, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	
Norcross, Jim	Champaign	IL	US	
Strand, David	Sherborn	MA	US	
Sweedler, Jonathan	Urbana	IL	US	

US-CL-CURRENT: 324/321; 324/306, 435/4

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	WWW	Draw De
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☐ 2. Document ID: US 20020130661 A1

L43: Entry 2 of 7

File: PGPB

Sep 19, 2002

PGPUB-DOCUMENT-NUMBER: 20020130661

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020130661 A1

TITLE: Nuclear magnetic resonance analysis of multiple samples

PUBLICATION-DATE: September 19, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Raftery, Daniel	Lafayette	IN	US	
Fisher, George G.	Oak Harbor	WA	US	

Petucci, Christopher J.	Memphis	TN	US
McNamara, Ernesto	Alexandria	VA	US

US-CL-CURRENT: 324/318; 324/309, 324/321, 324/322

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draws
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☐ 3. Document ID: US 20020105327 A1

L43: Entry 3 of 7

File: PGPB

Aug 8, 2002

PGPUB-DOCUMENT-NUMBER: 20020105327  
PGPUB-FILING-TYPE: new  
DOCUMENT-IDENTIFIER: US 20020105327 A1

TITLE: Steep solvent gradient NMR analysis method

PUBLICATION-DATE: August 8, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	
Norcross, Jim	Champaign	IL	US	

US-CL-CURRENT: 324/306; 324/321

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draws
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☐ 4. Document ID: US 6700379 B2

L43: Entry 4 of 7

File: USPT

Mar 2, 2004

US-PAT-NO: 6700379  
DOCUMENT-IDENTIFIER: US 6700379 B2

TITLE: Steep solvent gradient NMR analysis method

DATE-ISSUED: March 2, 2004

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Peck; Tim L.	Mahomet	IL		
Olson; Dean	Champaign	IL		
Norcross; Jim	Champaign	IL		

US-CL-CURRENT: 324/321; 324/306

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draws
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☐ 5. Document ID: US 6696838 B2

L43: Entry 5 of 7

File: USPT

Feb 24, 2004

US-PAT-NO: 6696838

DOCUMENT-IDENTIFIER: US 6696838 B2

TITLE: Nuclear magnetic resonance analysis of multiple samples

DATE-ISSUED: February 24, 2004

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Rafferty; Daniel	Lafayette	IN		
Fisher; George G.	Oak Harbor	WA		
McNamara; Ernesto	Alexandria	VA		

US-CL-CURRENT: 324/321; 324/310, 324/318, 324/322

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWIC	Draw De
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☐ 6. Document ID: US 6194900 B1

L43: Entry 6 of 7

File: USPT

Feb 27, 2001

US-PAT-NO: 6194900

DOCUMENT-IDENTIFIER: US 6194900 B1

TITLE: Integrated miniaturized device for processing and NMR detection of liquid phase samples

DATE-ISSUED: February 27, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Freeman; Dominique M.	Pescadero	CA		
Swedberg; Sally A.	Palo Alto	CA		

US-CL-CURRENT: 324/321; 324/318

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWIC	Draw De
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☐ 7. Document ID: US 6097188 A

L43: Entry 7 of 7

File: USPT

Aug 1, 2000

US-PAT-NO: 6097188

DOCUMENT-IDENTIFIER: US 6097188 A

TITLE: Microcoil based micro-NMR spectrometer and method

DATE-ISSUED: August 1, 2000

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Sweedler; Jonathan V.	Urbana	IL		
Magin; Richard L.	Urbana	IL		
Peck; Timothy L.	Champaign	IL		
Webb; Andrew G.	Urbana	IL		

US-CL-CURRENT: 324/321; 324/318, 324/322

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWC	Draw. Ds
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Term	Documents
RECEPTION	405653
RECEPTIONS	2969
DETECT\$6	0
DETECT	1169555
DETECTA	16
DETECTAB	1
DETECTABALE	9
DETECTABCLE	1
DETECTABE	9
DETECTABED	1
DETECTABEL	10
(L42 AND (DETECT\$6 OR RECEPTION OR RECEIV\$5 OR SENS\$5)).USPT,PGPB,JPAB,EPAB,DWPI,TDBD.	7

There are more results than shown above. [Click here to view the entire set.](#)

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Search Results - Record(s) 1 through 6 of 6 returned.

☐ 1. Document ID: US 20020149369 A1

Using default format because multiple data bases are involved.

L45: Entry 1 of 6

File: PGPB

Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020149369

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020149369 A1

TITLE: Microfluidic device with multiple microcoil NMR detectors

PUBLICATION-DATE: October 17, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	
Norcross, Jim	Champaign	IL	US	
Strand, David	Sherborn	MA	US	
Sweedler, Jonathan	Urbana	IL	US	

US-CL-CURRENT: 324/321; 324/306, 435/4

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Drawings
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☐ 2. Document ID: US 20020130661 A1

L45: Entry 2 of 6

File: PGPB

Sep 19, 2002

PGPUB-DOCUMENT-NUMBER: 20020130661

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020130661 A1

TITLE: Nuclear magnetic resonance analysis of multiple samples

PUBLICATION-DATE: September 19, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Raftery, Daniel	Lafayette	IN	US	
Fisher, George G.	Oak Harbor	WA	US	

Petucci, Christopher J.                      Memphis                      TN                      US  
McNamara, Ernesto                      Alexandria                      VA                      US

US-CL-CURRENT: 324/318; 324/309, 324/321, 324/322

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw. D.
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☐ 3. Document ID: US 20020105327 A1

L45: Entry 3 of 6

File: PGPB

Aug 8, 2002

PGPUB-DOCUMENT-NUMBER: 20020105327

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020105327 A1

TITLE: Steep solvent gradient NMR analysis method

PUBLICATION-DATE: August 8, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	
Norcross, Jim	Champaign	IL	US	

US-CL-CURRENT: 324/306; 324/321

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw. D.
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☐ 4. Document ID: US 6700379 B2

L45: Entry 4 of 6

File: USPT

Mar 2, 2004

US-PAT-NO: 6700379

DOCUMENT-IDENTIFIER: US 6700379 B2

TITLE: Steep solvent gradient NMR analysis method

DATE-ISSUED: March 2, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Peck; Tim L.	Mahomet	IL		
Olson; Dean	Champaign	IL		
Norcross; Jim	Champaign	IL		

US-CL-CURRENT: 324/321; 324/306

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw. D.
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☐ 5. Document ID: US 6696838 B2

L45: Entry 5 of 6

File: USPT

Feb 24, 2004

US-PAT-NO: 6696838

DOCUMENT-IDENTIFIER: US 6696838 B2

TITLE: Nuclear magnetic resonance analysis of multiple samples

DATE-ISSUED: February 24, 2004

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Rafferty; Daniel	Lafayette	IN		
Fisher; George G.	Oak Harbor	WA		
McNamara; Ernesto	Alexandria	VA		

US-CL-CURRENT: 324/321; 324/310, 324/318, 324/322

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWC	Draw Dc
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☐ 6. Document ID: US 6194900 B1

L45: Entry 6 of 6

File: USPT

Feb 27, 2001

US-PAT-NO: 6194900

DOCUMENT-IDENTIFIER: US 6194900 B1

TITLE: Integrated miniaturized device for processing and NMR detection of liquid phase samples

DATE-ISSUED: February 27, 2001

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Freeman; Dominique M.	Pescadero	CA		
Swedberg; Sally A.	Palo Alto	CA		

US-CL-CURRENT: 324/321; 324/318

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWC	Draw Dc
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Clear	Generate Collection	Print	Fwd Refs	Bkwd Refs	Generate OACS
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Term	Documents
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FUSED S	0
SILICA	436960
SILICA S	16978
PEEK	9980
PEEK S	322
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Search Results - Record(s) 1 through 4 of 4 returned.

☐ 1. Document ID: US 6696838 B2

Using default format because multiple data bases are involved.

L47: Entry 1 of 4

File: USPT

Feb 24, 2004

US-PAT-NO: 6696838

DOCUMENT-IDENTIFIER: US 6696838 B2

TITLE: Nuclear magnetic resonance analysis of multiple samples

DATE-ISSUED: February 24, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Raftery; Daniel	Lafayette	IN		
Fisher; George G.	Oak Harbor	WA		
McNamara; Ernesto	Alexandria	VA		

US-CL-CURRENT: [324/321](#); [324/310](#), [324/318](#), [324/322](#)

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	WMC	Draw Ds
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☐ 2. Document ID: US [6504368](#) B2

L47: Entry 2 of 4

File: USPT

Jan 7, 2003

US-PAT-NO: [6504368](#)

DOCUMENT-IDENTIFIER: US [6504368](#) B2

TITLE: Spectroscopic measurement method using NMR

DATE-ISSUED: January 7, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Ross; Alfred	Lorrach			DE
Schlotterbeck; Gotz	Efringen-Kirchen			DE
Senn; Hans	Windisch			CH

US-CL-CURRENT: [324/307](#); [324/308](#), [324/309](#), [324/318](#)

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KMC	Draw. De
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☐ 3. Document ID: JP 2004501350 W, WO 200179874 A1, EP 1158307 A1, US 20010045831 A1, AU 200173981 A, EP 1275011 A1, US 6504368 B2

L47: Entry 3 of 4

File: DWPI

Jan 15, 2004

DERWENT-ACC-NO: 2002-082759

DERWENT-WEEK: 200410

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TITLE: NMR spectrometer throughput increase using simultaneous determination of NMR spectra of samples arranged in a measuring site and varying the magnetic fields which reduces the sample volume required

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KMC	Draw. De
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☐ 4. Document ID: DE 4100915 A, KR 189041 B1, WO 9213270 A1, AU 9190514 A, DE 4100915 C2, EP 567464 A1, JP 06504368 W, US 5351029 A, EP 567464 B1, DE 59108090 G, ES 2091454 T3, JP 3053865 B2

L47: Entry 4 of 4

File: DWPI

Jul 16, 1992

DERWENT-ACC-NO: 1992-242890

DERWENT-WEEK: 200055

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TITLE: Carbon mon:oxide sensor used in gas mixt. contg. oxygen@ - comprises metal oxide doped with different metal oxide and catalytic metal oxide with high sensitivity

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KMC	Draw. De
------	-------	----------	-------	--------	----------------	------	-----------	--	--	--------	-----	----------

Clear	Generate Collection	Print	Fwd Refs	Bkwd Refs	Generate OACS
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Term	Documents
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"6504368".USPT,PGPB,JPAB,EPAB,DWPI,TDBD.	4
(6504368).USPT,PGPB,JPAB,EPAB,DWPI,TDBD.	4

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[Generate OACS](#)

Search Results - Record(s) 1 through 3 of 3 returned.

☐ 1. Document ID: US 20020149369 A1

Using default format because multiple data bases are involved.

L48: Entry 1 of 3

File: PGPB

Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020149369

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020149369 A1

TITLE: Microfluidic device with multiple microcoil NMR detectors

PUBLICATION-DATE: October 17, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	
Norcross, Jim	Champaign	IL	US	
Strand, David	Sherborn	MA	US	
Sweedler, Jonathan	Urbana	IL	US	

US-CL-CURRENT: 324/321; 324/306, 435/4

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	NMC	Draw De
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☐ 2. Document ID: US 20020105327 A1

L48: Entry 2 of 3

File: PGPB

Aug 8, 2002

PGPUB-DOCUMENT-NUMBER: 20020105327

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020105327 A1

TITLE: Steep solvent gradient NMR analysis method

PUBLICATION-DATE: August 8, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	

Norcross, Jim

Champaign

IL

US

US-CL-CURRENT: 324/306; 324/321

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	WMC	Draw De
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☐ 3. Document ID: US 6700379 B2

L48: Entry 3 of 3

File: USPT

Mar 2, 2004

US-PAT-NO: 6700379

DOCUMENT-IDENTIFIER: US 6700379 B2

TITLE: Steep solvent gradient NMR analysis method

DATE-ISSUED: March 2, 2004

## INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Peck; Tim L.	Mahomet	IL		
Olson; Dean	Champaign	IL		
Norcross; Jim	Champaign	IL		

US-CL-CURRENT: 324/321; 324/306

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	WMC	Draw De
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Clear	Generate Collection	Print	Fwd Refs	Bkwd Refs	Generate OACS
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Term	Documents
FIELD	3201991
FIELDS	453756
GRADIENT	246992
GRADIENTS	60840
DYNAMIC\$4	0
DYNAMIC	468225
DYNAMICA	33
DYNAMICAAY	2
DYNAMICABLY	1
DYNAMICABY	12
DYNAMICADY	6
(L43 AND ((DYNAMIC\$4 OR ELECTRIC\$4) WITH FIELD WITH GRADIENT WITH FOCUS\$4)).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	3



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L48: Entry 2 of 3

File: PGPB

Aug 8, 2002

PGPUB-DOCUMENT-NUMBER: 20020105327

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020105327 A1

TITLE: Steep solvent gradient NMR analysis method

PUBLICATION-DATE: August 8, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	
Norcross, Jim	Champaign	IL	US	

## ASSIGNEE-INFORMATION:

NAME	CITY	STATE	COUNTRY	TYPE CODE
Protasis Corporation	Marborough	MA	US	02

APPL-NO: 10/ 006503 [PALM]

DATE FILED: December 3, 2001

## RELATED-US-APPL-DATA:

Application is a non-provisional-of-provisional application 60/250705, filed December 1, 2000,

INT-CL: [07] G01 V 3/00

US-CL-PUBLISHED: 324/306; 324/321

US-CL-CURRENT: 324/306; 324/321

## ABSTRACT:

An NMR method of analyzing an analyte comprises feeding an analyte sample fluid to an NMR flow cell. The NMR flow cell comprises an RF microcoil operably associated with an enlarged containment region. The mobile phase of the analyte sample flowing through the NMR flow cell has a solvent gradient greater than 10% per minute. The analyte sample fluid can be fed to the NMR flow cell from an analyte extraction chamber, e.g., operative to perform liquid chromatography, capillary electrophoresis, or the like, especially a capillary-based analyte extraction chamber integrated in an NMR probe with the NMR flow cell. A sample volume is held in the NMR flow cell for equilibration less than 1 hour, preferably less than 30 minutes prior to actuating NMR analysis of the observe volume in the microcoil.

# Hit List

Search Results - Record(s) 1 through 1 of 1 returned.

☐ 1. Document ID: US 20020149369 A1

Using default format because multiple data bases are involved.

L50: Entry 1 of 1

File: PGPB

Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020149369

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020149369 A1

TITLE: Microfluidic device with multiple microcoil NMR detectors

PUBLICATION-DATE: October 17, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Peck, Tim L.	Mahomet	IL	US	
Olson, Dean	Champaign	IL	US	
Norcross, Jim	Champaign	IL	US	
Strand, David	Sherborn	MA	US	
Sweedler, Jonathan	Urbana	IL	US	

US-CL-CURRENT: 324/321; 324/306, 435/4

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Drawings
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Term	Documents
FLUID	1663992
FLUIDS	351052
ROUT\$4	0
ROUT	5226
ROUTA	32
ROUTAB	1
ROUTABLE	1038
ROUTABLY	12

ROUTACT	3
ROUTACTS	1
ROUTAC4	1
(L39 AND (FLUID WITH ROUT\$4)).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	1

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FILE 'INPADOC, WPIX, HCAPLUS, JAPIO' ENTERED AT 14:18:50 ON 01 JUN 2004

E US2000-0250874/PRN,AP

L1 5 S (US2000-250874P/PRN OR US2000-250874P/AP)

FILE 'DPCI' ENTERED AT 14:20:03 ON 01 JUN 2004

E US2000-0250874/PRN,AP

E WO 2002056049/PN

FILE 'SCISEARCH' ENTERED AT 14:36:58 ON 01 JUN 2004

E LI Y, 1999, V71/RE

L2 19 S "LI Y, 1999, V71, P4815, ANAL CHEM"/RE

L3 0 S L2 AND (PY=1999 OR PY=20000)

L4 1 S L2 AND (PY=1999 OR PY=2000)

*NPC STIC Search*  
*Jan 1<sup>st</sup> 2004*  
*See attached Search History Database*  
*9 Records TAF*

FILE 'INPADOC, WPIX, HCAPLUS, JAPIO' ENTERED AT 14:40:52 ON 01 JUN 2004

L5 162 S PECK?/AU,IN AND (REFOCUS? OR FOCUS?)

L6 1 S L5 AND DYNAMIC?

L7 2 S US6696838/PN

FILE 'HCAPLUS' ENTERED AT 14:44:32 ON 01 JUN 2004

L8 3935 S (FOCUS##### OR REFOCUS#####)(3A)(ELECTR  
IC### OR FIELD OR DYNAMIC#####)L9 38890 S MICROFLUID##### OR MICROCOIL##### OR  
MICROPROB##### OR MICROCOIL##### OR MICRO###(2A)(COIL OR PROBE OR COIL OR FLUID###)L10 28009 S (MULTI OR MULTIPLE OR SEVERAL)(3A)(PATH####  
## OR CHANNEL### OR CONDUIT) OR MULTICHANNEL? OR MULTIPATH? OR MULTICONDUIT?

L11 0 S L8 AND L9 AND L10

L12 27 S L8 AND L9

L13 21 S L8 AND L10

L14 1227 S (REFOCUS? OR FOCUS?) AND L9

L15 14 S L14 AND L10

L16 18 S ((L12 OR L13) OR L15) AND (ROUT##### OR  
SWITCH##### OR DIRECT##### OR MICROROUT? OR MICROSWITCH? OR MICRODIRECT?)

L17 10 S L16 AND SAMPL#####

L18 0 S L16 AND (NMR OR MRI OR MR OR MAGNETIC RESONANCE OR NUCLEAR MAGNETIC)

L19 672 S (L8 OR L9 OR L10) AND (NMR OR MRI OR MR OR MAGNETIC RESONANCE OR NUCLEAR  
MAGNETIC)

L20 52 S L19 AND DYNAMIC#####

L21 233 S L19 AND MULTI#####

L22 19 S L20 AND L21

L23 81 S (L12 OR L13) OR (L15 OR L16 OR L17) OR L22

L24 7 S L23 AND P/DT

L25 21 S L23 AND (CONTROL##### OR MICROCONTROL#### #####)

L26 18 S L25 NOT L24

L27 7 S L26 NOT PY&gt;2000

L28 6 S L24 NOT PRY&gt;2000

L29 14 S L27 OR L28 OR L24

FILE 'HCAPLUS' ENTERED AT 14:58:23 ON 01 JUN 2004

L30 420322 S G01R033?/IC OR G01V003?/IC OR NMR OR MRI OR MAGNETIC RESONANCE

L31 552 S (L12 OR L13 OR L14 OR L15 OR L16 OR L17 OR  
L18 OR L19 OR L20 OR L21 OR L22 OR L23 OR L24 OR L25) AND L30

L32 69 S L31 AND (FOCUS##### OR REFOCUS#####)

L33 48 S L31 AND DYNAMIC#####

L34 7 S L31 AND ELECTRIC## FIELD

FILE 'HCAPLUS' ENTERED AT 14:58:23 ON 01 JUN 2004

L35 137 S L31 AND (CHANNELS OR PATHS OR PATHWAYS OR CONDUITS)  
 L36 1 S L31 AND (MICROCHANNELS OR MICROPATHS OR MICROPATHWAYS OR MICROCONDUITS)  
 L37 172 S L31 AND MULTI#####  
 L38 0 S L31 AND MANIFOLD  
 L39 27 S L32 AND L33  
 L40 3 S L32 AND L34  
 L41 1 S L33 AND L34  
 L42 18 S (L32 OR L33 OR L34) AND L35  
 L43 1 S (L32 OR L33 OR L34) AND L36  
 L44 24 S (L32 OR L33 OR L34) AND L37  
 L45 259 S L31 AND (MICRO#####(2A)(COIL OR PROBE OR FLUID#####) OR MICROCOIL? OR  
 MICROPROB? OR MICROFLUID?)  
 L46 7 S (L39 OR L40 OR L41 OR L42 OR L43 OR L44) AND L45  
 L47 13 S L34 OR L36 OR (L40 OR L41) OR L43 OR L46

FILE 'SCISEARCH' ENTERED AT 15:06:55 ON 01 JUN 2004

E WU N, 1994, V66/RE

L48 3 S ("WU N, 1994, V66, P384, ANAL CHEM"/RE OR "WU N, 1994, V66, P3894, ANAL CHEM"/RE)

FILE 'ANABSTR' ENTERED AT 15:07:44 ON 01 JUN 2004

L49 0 S CHARACTERISATION AND SUBSTANCES SEPARATED AND CHROMATOGRAPHY  
 L50 0 S CHARACTERI##### AND SUBSTANCES SEPARATED AND CHROMATOGRAPHY

FILE 'BIOSIS' ENTERED AT 15:08:20 ON 01 JUN 2004

L51 2 S CHARACTERI##### AND SUBSTANCES SEPARATED AND CHROMATOGRAPHY

FILE 'HCAPLUS' ENTERED AT 15:09:00 ON 01 JUN 2004

L52 248 S COILS(L)MICRO  
 L53 3 S L52 AND MICROCHANNEL?  
 L54 14 S NMR AND L52

FILE 'ANABSTR' ENTERED AT 15:13:56 ON 01 JUN 2004

L55 5016 S ("NUCLEAR MAGNETIC RESONANCE"/CT OR  
 "NUCLEAR MAGNETIC RESONANCE (NMR)"/CT) OR NMR  
 E MICROCOIL? OR MICROPROB? OR MICROFLUID?  
 L56 1067 S MICROCOIL? OR MICROPROB? OR MICROFLUID?  
 L57 1641 S MICRO COIL? OR MICRO PROB? OR MICRO FLUID?  
 L58 1451 S (MULTI OR MULTIPLE OR FOUR OR FIVE OR SIX  
 IR SEVEN OR EIGHT OR NINE OR TEN OR TWO OR THREE)(2W)(PATH#####  
 ## OR CHANNEL##### OR CONDUIT##### OR ROUT##### OR DIRECTION#####)  
 L59 377 S L55 AND MULTI#####  
 L60 65 S L55 AND DYNAMIC#####  
 L61 256 S L55 AND FIELD  
 L62 26 S L55 AND ELECTRIC#####  
 L63 46 S L59 AND (L60 OR L61 OR L62)  
 L64 32 S L55 AND (L56 OR L57 OR L58)  
 L65 2 S L63 AND L64  
 L66 52 S L59 AND (L60 OR L61 OR L62 OR L63 OR L64 OR L65)  
 L67 25 S L66 AND PY>2000  
 L68 27 S L66 NOT L67  
 L69 46 S PECK?/AU  
 L70 3 S L55 AND L69  
 L71 9 S L68 AND (MU OR MUM OR MICRO#####)

FILE 'ANABSTR, BIOSIS, HCAPLUS, INSPEC' ENTERED AT 15:23:36 ON 01 JUN 2004

L72 482 S HANER?/AU,IN OR DECHOW?/AU,IN  
L73 26 S L72 AND (MICRODIRECT? OR MICROROUT? OR  
ROUT##### OR DIRECTION##### OR SWITCH#####)  
L74 56 S L72 AND (MICROCONTROL? OR CONTROL?)  
L75 3 S (L73 OR L74) AND (MR OR NMR OR MAGNETIC RESONANCE)  
L76 64884 S FLOWTHROUGH OR FLOW THROUGH OR SAMPLE DELIVERY  
L77 605 S L76 AND (NMR OR MR OR MAGNETIC RESONANCE)  
L78 72 S L77 AND MULTI#####  
L79 15 S L78 AND MICRO#####

L47 ANSWER 8 OF 13 HCAPLUS COPYRIGHT 2004 ACS on STN  
AN 1995:449152 HCAPLUS  
DN 123:186477  
ED Entered STN: 29 Mar 1995  
TI Fast **multichannel** stabilization of the **magnetic resonance** in a magneto-resonance spectrometer  
AU Borisov, Yu. V.; Ivanov, S. N.; Lobashev, V. M.; Sobolev, Yu. V.  
CS Petersburg Nuclear Physics Institute (PNPI), Gatchina, Leningrad district, 188350, Russia  
SO Nuclear Instruments & Methods in Physics Research, Section A: Accelerators, Spectrometers, Detectors, and Associated Equipment (1995), 357(1), 115-19  
CODEN: NIMAER; ISSN: 0168-9002  
PB Elsevier  
DT Journal  
LA English  
CC 77-7 (Magnetic Phenomena)  
AB A method of fast stabilization of the **magnetic resonance** in a **magnetic resonance** spectrometer was developed and tested. The source of the oscillatory field is the synthesizer which frequency follows variations of the average magnetic field in spectrometer. To search for the neutron elec. dipole moment in the exptl. setup under unfavorable conditions in the reactor exptl. hall a stability of the **magnetic resonance** was achieved that is equivalent to a magnetic field stability at the level of  $\approx 2.5 \times 10^{-12}$  T per 6 min in a volume of  $\approx 50$  L with an **elec. field** of 1.5 MV/m.

N/A TAF 6/1/2004



L47 ANSWER 11 OF 13 HCAPLUS COPYRIGHT 2004 ACS on STN  
 AN 1976:569155 HCAPLUS  
 DN 85:169155  
 ED Entered STN: 12 May 1984  
 TI Tensor polarizability of the ground-state hyperfine structure of thallium  
 AU Gould, Harvey  
 CS Dep. Phys., Brandeis Univ., Waltham, MA, USA  
 SO Physical Review A: Atomic, Molecular, and Optical Physics (1976), 14(3),  
 922-7  
 CODEN: PLRAAN; ISSN: 1050-2947  
 DT Journal  
 LA English  
 CC 73-3 (Spectra by Absorption, Emission, Reflection, or Magnetic Resonance,  
 and Other Optical Properties)  
 AB The atomic-beam **magnetic-resonance** technique was used to  
 measure the hyperfine-structure tensor polarizability (quadratic Stark  
 effec) in the 62P<sub>1/2</sub> ground state of atomic Tl. **Elec.**  
**fields** of up to 460 kV/cm were used to lift the degeneracy between  
 the  $m_F = 0$  and the  $m_F = \pm 1$  substrates in the absence of an external  
 magnetic **field**, and **focusing** transitions between these  
 Stark-separated states were observed. Measurements were also made between  
 Zeeman-separated substates. The results are  $aT = -(3.74 \pm 0.09) +$   
 $10^{-8} \text{ Hz}/(\text{V}/\text{cm})^2$ , or  $k = -(5.62 \pm 0.14) + 10^{-8} \text{ Hz}/(\text{V}/\text{cm})^2$ , where  
 $\delta v = kE^2$  is the Stark shift of the ( $m_F = 0$ ) .dblarw. ( $m_F = -1$ )  
 flop-in transition.

N/A TAF 6/1/2004

L47 ANSWER 12 OF 13 HCAPLUS COPYRIGHT 2004 ACS on STN  
AN 1968:108935 HCAPLUS  
DN 68:108935  
ED Entered STN: 12 May 1984  
TI Test methods used in research on insulating materials  
AU Stamm, Hans  
CS Tech. Hochsch., Ilmenau, Fed. Rep. Ger.  
SO Elektr. (1967), 21(11), 411-13  
CODEN: EKTRAO; ISSN: 0013-5399  
DT Journal  
LA German  
CC 71 (Electric Phenomena)  
AB Methods for determining fatigue in insulators are surveyed. Reversible and irreversible fatigue are distinguished and their dependence upon mol. and crystal structure is discussed. The usual method for the determination of fatigue is the d.c. charging and discharging as a function of time and temperature. The interpretation of the obtained curve is difficult and no prediction about breakdown is possible. A direct relation between structural changes and dielec. consts. has been found for crystals. The relevant structure parameters are dipoles (mobility and orientation) and ions (mobility) and their dependence upon **elec. fields**. The dependence of the absorption spectra on temperature gives information about (1) the mobility and orientation of dipoles and (2) the breaking of chains and/or changes of configuration which are also related to changes of viscosity. Information regarding changes of elec. dipoles is obtained from ir spectra and changes of polarizability from the Raman spectra. In order to obtain a representative description of structure and structural changes, other analyzing methods must be introduced. Preferable are nondestructive methods and the following are considered: microscopic methods for the study of texture, domains due to **elec. fields**, and the followup of quick changes in dielects. by the Kerr effect with polarized light. Order-disorder phenomena and dislocations can be observed by electron microscopy by using e.g. deflection of the electron beam by magnetic or **elec. fields**. Studies at 100-300° can be informative. The configuration of electron shells of specific atoms, charge and spin ds., and the structure and symmetry of the lattice on an atomic scale can be obtained by neutron diffraction and **magnetic resonance** methods. Structural details of the surface, especially after ion etching, can be observed by emission electron microscopy. The importance of the preceding methods is their information about stacking faults, which influence strongly the elec. properties of insulators. The orientation of the crystal axes relative to the geometric shape can be determined by diffraction methods, e.g. grazing incidence electrons or x-ray Kikuchi lines. Accurate detns. of lattice consts. can give information about impurities in the lattice. Constituent elements in small surface areas can be determined by the **microprobe** technique. By thermogravimetric measurements the actual aging of insulators could be studied by short time tests.

N/A TAF 6/1/2004

L47 ANSWER 13 OF 13 HCAPLUS COPYRIGHT 2004 ACS on STN  
AN 1962:447154 HCAPLUS  
DN 57:47154  
OREF 57:9372h-i,9373a-b  
ED Entered STN: 22 Apr 2001  
TI Radiospectroscopy: a new branch of modern physics  
AU Pekarek, Ludek  
SO Pokroky Mat. Fys. Astron. (1959), 4, 42-53;162-79  
DT Journal  
LA Unavailable  
CC 10 (Spectra and Some Other Optical Properties)  
AB Radiospectroscopy deals with absorption in the mm. and cm. range, and is divided into high-frequency spectra of gases and **magnetic resonance** spectra of solids and liquids. A high-frequency spectrograph is described. Spectra of NH<sub>3</sub>, SCSe, and HCN are shown. The absorption lines are attributed to mol. rotation and bending. The spectral lines, which are very narrow at low temps. and pressures, are widened at room temperature and atmospheric pressure. Two types of atomic clocks, both using the 23,870.11Mc. line of NH<sub>3</sub>, but one using a klystron source with its frequency regulated by the absorption by NH<sub>3</sub> mols., the other using excited NH<sub>3</sub> mols. themselves (separated from those in the ground state by **focusing** in an inhomogeneous **elec. field**) as sources. A region of overlap of microwave and infrared spectroscopy (wavelengths 0.7-1 mm.) is discussed; the former is more accurate. The nature of electron paramagnetic resonance and its phys. mechanism is explained. Applications in determining the structure of solid crystals, measuring magnetic fields, and following chemical reactions involving organic radicals are briefly outlined. A similar discussion is given of **nuclear magnetic resonance**, with emphasis on the very sharp peaks observed, and applications in accurate measurement of magnetic fields, and for their stabilization. Chemical fine structure and its use in chemical analysis is mentioned.

NA TAF 6/1/2004

L54 ANSWER 8 OF 14 HCAPLUS COPYRIGHT 2004 ACS on STN  
AN 2000:778333 HCAPLUS  
ED Entered STN: 07 Nov 2000  
TI Triaxial magnetic field gradient system for microcoil magnetic resonance  
imaging  
AU Seeber, D. A.; Hoftiezer, J. H.; Daniel, W. B.; Rutgers, M. A.;  
Pennington, C. H.  
CS Department of Physics, The Ohio State University, Columbus, OH, 43210, USA  
SO Review of Scientific Instruments (2000), 71(11), 4263-4272  
CODEN: RSINAK; ISSN: 0034-6748  
PB American Institute of Physics  
DT Journal  
LA English  
AB There is a great advantage in signal to noise ratio (S/N) that can be  
obtained in **NMR** (**NMR**) expts. on very small samples  
(having spatial dimensions .apprx.100  $\mu\text{m}$  or less) if one employs  
**NMR "micro" receiver coils, "microcoils,"**  
which are of similarly small dimensions. The gains in S/N could enable  
magnetic resonance imaging (MRI) microscopy with spatial resolution of  
.apprx.1-2  $\mu\text{m}$ , much better than currently available. Such MRI  
microscopy however requires very strong (>10 T/m), rapidly switchable  
triaxial magnetic field gradients. Here, we report the design and  
construction of such a triaxial gradient system, producing gradients  
substantially greater than 15 T/m in all three directions, x, y, and z  
(and as high as 50 T/m for the x direction). The gradients are switchable  
within time .apprx.10  $\mu\text{s}$  and adequately uniform (within 5% over a volume  
of [600 $\mu\text{m}$ 3] for microcoil MRI of small samples.).

N/A TAF 6/1/2004

L54 ANSWER 9 OF 14 HCAPLUS COPYRIGHT 2004 ACS on STN  
AN 2000:598842 HCAPLUS  
DN 133:260236  
ED Entered STN: 29 Aug 2000  
TI Fabrication of **NMR**-microsensors for nanoliter sample volumes  
AU Dechow, J.; Forchel, A.; Lanz, T.; Haase, A.  
CS Technische Physik, Universitat Wurzburg, Wurzburg, D-97074, Germany  
SO Microelectronic Engineering (2000), 53(1-4), 517-519  
CODEN: MIENEF; ISSN: 0167-9317  
PB Elsevier Science B.V.  
DT Journal  
LA English  
CC 76-14 (Electric Phenomena)  
AB The fabrication of **micro**-sensors for **NMR** spectroscopy on both glass and GaAs is presented. Planar **coils** with inner diameter from 50  $\mu\text{m}$  to 400  $\mu\text{m}$  including a coplanar wave-guide leading to the bonding pads were combined with a chamber for liquid samples of 200-500  $\mu\text{m}$  diameter on the backside of the substrate. The microcoil served as a receiver in a **1H-NMR** experiment at 11T (500 MHz). In initial expts., the spectrum of 60 nl-vols. of pure silicone-oil were detected by the microcoil.

NA TAF 6/1/2004

L54 ANSWER 10 OF 14 HCAPLUS COPYRIGHT 2004 ACS on STN  
AN 2000:45466 HCAPLUS  
DN 132:200809  
ED Entered STN: 19 Jan 2000  
TI Development and characterization of an **NMR** microsensor for  
nanoliter sample volumes  
AU Dechow, Joern; Forchel, Alfred W. B.; Lanz, Titus; Haase, Axel  
CS Tech. Phys., Univ. Wuerzburg, Wuerzburg, Germany  
SO Proceedings of SPIE-The International Society for Optical Engineering  
(1999), 3857 (Chemical Microsensors and Applications II), 98-103  
CODEN: PSISDG; ISSN: 0277-786X  
PB SPIE-The International Society for Optical Engineering  
DT Journal  
LA English  
CC 73-11 (Optical, Electron, and Mass Spectroscopy and Other Related  
Properties)  
AB The fabrication and performance of a **micro**-sensor for  
**NMR**- spectroscopy of nanoliter-sample vols. is presented. On both  
glass and GaAs-substrate, planar **coils** with inner diameter at  
50-400  $\mu\text{m}$  including a coplanar wave-guide leading to the bonding pads  
were fabricated. A chamber for the liquid samples of 200-500  $\mu\text{m}$  diameter  
was etched isotropically on the backside of the substrate, located under  
the coil. In initial expts., the spectrum of a 20-50 nL-vols. of pure  
Si-oil is analyzed in a 1H-**NMR** experiment in a 11T spectrometer (500  
MHz). The microcoil serves as a receiver, while the RF-power was  
transmitted by a macroscopic coil perpendicular to the receiver coil. The  
authors observe characteristic lines from the Si-oil spectrum which  
clearly indicates the high sensitivity of the microcoil. Addnl. signal  
from different materials in the experiment are suppressed by gradient fields  
and an adequate design of the sensor.

NA TAF 6/1/2004

L54 ANSWER 12 OF 14 HCAPLUS COPYRIGHT 2004 ACS on STN  
AN 1999:585088 HCAPLUS  
DN 131:306163  
ED Entered STN: 20 Sep 1999  
TI High-Resolution **NMR** Spectroscopy of Sample Volumes from 1 nL to  
10  $\mu$ L  
AU Lacey, Michael E.; Subramanian, Raju; Olson, Dean L.; Webb, Andrew G.;  
Sweedler, Jonathan V.  
CS Department of Chemistry Department of Electrical and Computer Engineering  
and the Beckman Institute, University of Illinois at Urbana-Champaign,  
Urbana, IL, 61801, USA  
SO Chemical Reviews (Washington, D. C.) (1999), 99(10), 3133-3152  
CODEN: CHREAY; ISSN: 0009-2665  
PB American Chemical Society  
DT Journal; General Review  
LA English  
CC 77-0 (Magnetic Phenomena)  
Section cross-reference(s): 80  
AB A review with 143 refs. Through the fabrication of nanoliter-volume  
**NMR** probes and their coupling to micro-separation strategies,  
mass-limited analytes in complex matrixes are becoming viable samples for  
**NMR** anal. Recent demonstrations of online HPLC-**NMR**-MS  
have combined one of the most widely used separation methods with two of the  
most information-rich techniques of chemical characterization. The extension  
of this double hyphenated method to the capillary scale will enable rapid,  
chemical rich screening of mass-limited samples with enhanced mass  
sensitivity. While the increased sensitivity of reduced-diameter radio  
frequency (RF) probes provides a widely applicable benefit for **NMR**  
spectroscopy,, microcoils offer several addnl. advantages which have not  
been fully explored. As one example, the diminutive spatial dimensions of  
microcoil probes enable their use in low-homogeneity high-field magnets.  
Microcoils can also be applied to solid-state **NMR** and the use of  
reduced-diameter RF probes for micro-imaging and **NMR** spectroscopic  
characterization of cellular samples has only begun to receive attention.  
Numerous areas in the biol. sciences will undoubtedly benefit from  
improved **NMR** spectroscopy of small vols. in the coming decade.

NA TAF 6/1/2004

Already of Record by Applicant's INS

L70 ANSWER 1 OF 3 ANABSTR COPYRIGHT 2004 RSC on STN  
AN 58(2):F86 ANABSTR  
TI Online **NMR** detection of amino-acids and peptides in microbore  
LC.  
AU Wu, N.; Webb, A.; **Peck, T. L.**; Sweedler, J. V. (Beckman Inst.  
Adv. Sci. and Technol., Univ. Illinois, Urbana, IL 61801, USA)  
SO Anal. Chem. (1995) 67(18), 3101-3107  
CODEN: ANCHAM ISSN: 0003-2700  
DT Journal  
LA English  
AB A model was developed to predict signal-to-noise ratios in flow systems as  
a function of flow rate, sample volume, microcil size and **NMR**  
acquisition parameters. The model predicts that a reduction of detector  
volume from 16  $\mu$ l to 50 nl would decrease the signal-to-noise ratio by  
2. This was tested by the analysis of three amino-acids and two peptides  
on a column (15 cm + 1 mm i.d.) of C18 (5  $\mu$ m). The column was in  
the bore of a magnet and connected to a microcil (14-17 turns of varnished  
42-gauge Cu wire on a capillary [355  $\mu$ m o.d., 250  $\mu$ m i.d.]). Elution  
(10-50  $\mu$ l/min) was with deuterated acetonitrile/2% TFA in D2O (13:47).  
Scan number was 64, 128 or 256, requiring 9, 18 or 36 s with a pulse  
repetition rate of 0.03 or 0.06 s. These conditions allowed good  
**NMR** spectra to be obtained from 1  $\mu$ g of analyte in the 50 nl  
cell described.

NA TAF 6/1/2004



L71 ANSWER 2 OF 9 ANABSTR COPYRIGHT 2004 RSC on STN  
AN 62(11):F10171 ANABSTR  
TI A **microcoil NMR** probe for coupling **microscale**  
HPLC with on-line **NMR** spectroscopy.  
AU Subramanian, R.; Kelley, W. P.; Floyd, P. D.; Tan, Z. J.; Webb, A. G.;  
Sweedler, J. V. (jsweedle@uiuc.edu, Dept. Chem., Beckman Inst. Advanced  
Sci. and Technol., Univ. Illinois, Urbana, IL 61801, USA)  
SO Anal. Chem. (1999) 71(23), 5335-5339  
CODEN: ANCHAM ISSN: 0003-2700  
DT Journal  
LA English  
AB An HPLC **NMR** system is presented that integrates a commercial  
**microbore** HPLC system using a 0.5 mm column with a 500 MHz proton  
**NMR** spectrometer using a customer **NMR** probe with an  
observe volume of 1.1  $\mu\text{L}$  and a coil fill factor of 68%.  
Careful attention to capillary connections and **NMR** flow cell  
design allows on-line **NMR** detection with no significant loss in  
separation efficiency when compared with a UV chromatogram. HPLC  
**NMR** is performed on mixtures of amino acids and small peptides  
with analyte injection amounts as small as 750 ng; the separations are  
accomplished in less than 10 min and individual **NMR** spectra are  
acquired with 12 s time resolution. Stopped-flow **NMR** is achieved  
by diversion of the chromatographic flow after observation of the  
beginning of the analyte band within the **NMR** flow cell.  
Isolation of the compound of interest within the **NMR** detection  
cell allows **multidimensional** experiments to be performed. A  
stopped-flow COSY spectrum of the peptide Phe-Ala is acquired in 3.5 h  
with an injected amount of 5  $\mu\text{g}$ .

NA TAF 6/1/2004

L71 ANSWER 4 OF 9 ANABSTR COPYRIGHT 2004 RSC on STN  
 AN 61(1):F32 ANABSTR  
 TI Investigation of the calcium content in joint cartilage: is it connected  
 with (early arthrotic) changes in cartilage structure?  
 AU Reinert, T.; Butz, T.; Flagmeyer, R.-H.; Jankuhn, S.; Vogt, J.; Gruender,  
 W.; Kanowski, M.; Wagner, M.; Werner, A.; Grambole, D.; Herrmann, F.  
 (reinert@physik.uni-leipzig.de, Fac. Phys. and Geol., Univ. Leipzig, 04103  
 Leipzig, Germany)  
 SO Nucl. Instrum. Methods Phys. Res., Sect. B (1998) B136-B138, 936-940  
 CODEN: NIMBEU ISSN: 0168-583X  
 (Presented at the Thirteenth International Conference on Ion Beam  
 Analysis, held in Lisbon, Portugal, 27 Jul-1 Aug 1997)  
 DT Journal  
 LA English  
 AB Cylinders (diameter 1.8 cm) of porcine femoral cartilage and bone were  
 examined by **NMR** imaging, and a cylinder (diameter 3 mm)  
 extending from the articular surface to the bone was punched from those  
 that showed marked **multilaminar** structure. The cylinder was  
 sectioned transversely (200 **.mu.m**) to isolate tissue at  
 particular distances from the bone towards the articular surface, and the  
 sections were dried by centrifugation and mounted on Al for PIXE. The  
 sections were bombarded with 1.7 MeV protons, and the X-rays were detected  
 with an Si(Li) detector at 141.2° to the proton beam  
 (back-scattered protons detected with a PIPS detector at 170° to  
 the beam). For better lateral resolution, sections (200 **.mu.m**)  
 perpendicular to the articular surface were also cut to correspond to the  
**NMR** images, and the dried and mounted sections were bombarded with  
 a 3 MeV proton **micro-probe** beam focused to 20 **.mu.m**  
 in the scanning direction (fast-scanning mode) perpendicular  
 to the articular surface, the X-rays being detected with an Si(Li)  
 detector at 120° to the proton beam. Attempts to correlate the Ca  
 concentrations with the **NMR** images were not immediately  
 successful.

NA TAF 6/1/2004

L71 ANSWER 5 OF 9 ANABSTR COPYRIGHT 2004 RSC on STN  
AN 60(12):C39 ANABSTR  
TI Design of solenoidal **microcoils** for high-resolution carbon-13  
**NMR** spectroscopy.  
AU Subramanian, R.; Webb, A. G. (Dept. Electrical and Computer Eng. and  
Beckman Inst. Adv. Sci. and Technol., Univ. Illinois at Urbana-Champaign,  
Urbana, IL 61801, USA)  
SO Anal. Chem. (1998) 70(13), 2454-2458  
CODEN: ANCHAM ISSN: 0003-2700  
DT Journal  
LA English  
AB Two **microcoil** detection probes (circuit diagrams presented)  
suitable for smaller-sized samples than the mg amounts required hitherto  
have been developed to improve detection limits in high-resolution  
natural-abundance <sup>13</sup>C **NMR** at 11.7 T. Both probes have a  
coil-filling factor of 69%. The direct detection probe incorporates <sup>13</sup>C  
observe, <sup>2</sup>H lock and <sup>1</sup>H decouple channels and has an observe volume of 1 .  
**mu**.l. A linewidth of 1.3 Hz was achieved, and the detection limits  
for sucrose (natural abundance) and (3-<sup>13</sup>C)-L-alanine (99%) were 52 and  
4.2 nmol with acquisition times of 90 min and <30 s, respectively. The  
inverse detection probe incorporates <sup>1</sup>H observe and <sup>13</sup>C decouple channels  
and has an observe volume of 550 nl. The detection limit for sucrose from  
a 1D <sup>13</sup>C-decoupled heteronuclear **multiple**-quantum coherence  
spectrum was 4.5 nmol with an acquisition time of 14 min.

NA TAI<sup>2</sup> 6/1/2004

L71 ANSWER 6 OF 9 ANABSTR COPYRIGHT 2004 RSC on STN  
AN 60(6):H222 ANABSTR  
TI Liquid chromatography coupled to mass spectrometry and nuclear magnetic resonance spectroscopy for the screening of plant constituents.  
AU Wolfender, J.-L.; Rodriguez, S.; Hostettmann, K. (Inst. Pharmacognosie et Phytochim., Univ. Lausanne, 1015 Lausanne, Switzerland)  
SO J. Chromatogr., A (1998) 794(1-2), 299-316  
CODEN: JCRAEY ISSN: 0021-9673  
DT Journal  
LA English  
AB Coupling methods of HPLC with diode-array UV, single and tandem MS and **NMR** are assessed for the phytochemical screening of crude plant extracts. Compounds considered include xanthenes, their O-glycosides and aglycones, triterpene glycosides (saponins) and polyphenols. For HPLC, a column (15 cm + 3.9 mm i.d.) of Nova-Pak C18 (4  $\mu$ m) was used with a similarly packed guard column. For MS thermospray for electrospray interfaces were used and/or modes of continuous-flow FAB MS or tandem or **multiple**-cycle MS. For **NMR** a Unity Innova 500 MHz instrument with Varian  $^1\text{H}/^{13}\text{C}$  pulse **field**-gradient indirect-detection **microflow** LC-**NMR** probe (60- $\mu$ l flow cell, 3 mm o.d.) was used. A range of chromatograms and mass spectrograms derived from analyses of Gentianaceae and Leguminosae specimens are presented and results are discussed. Use of the diverse coupling techniques can eliminate the need for time-consuming analyte isolation procedures.

N/A TAF 6/1/2004

L75 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2004 ACS on STN  
 AN 1998:13756 HCAPLUS  
 DN 128:56837  
 ED Entered STN: 10 Jan 1998  
 TI Sample delivery system used in chemical analysis methods which employs  
 pressurized gas for sample conveyance  
 IN **Haner, Ronald L.**; Duff, David W.; Kellogg, Christopher C.  
 PA Varian Associates, Inc., USA  
 SO Eur. Pat. Appl., 20 pp.  
 CODEN: EPXXDW  
 DT Patent  
 LA English  
 IC ICM G01R033-30  
 ICS G01N001-10  
 CC 79-2 (Inorganic Analytical Chemistry)  
 Section cross-reference(s): 77, 80  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 813071	A1	19971217	EP 1997-304170	19970613
	JP 10090383	A2	19980410	JP 1997-171277	19970613
PRAI	US 1996-665165	A	19960614		

AB A sample delivery system for a flow-through **NMR** anal. is provided, which uses pressurized gas as a means for conveying a sample into and out of an **NMR** spectrometer. Two sources of gas pressure, a forward pressure and back pressure, oppose the sample within the tubing of the sample delivery system and the tubing of the flow-through system which are operatively coupled together. Conveyance of the sample in any **direction** within the tubing is achieved by adjusting the pressure differential. Precise positioning of the sample in the magnetic field center and complete removal of the sample from the **NMR** spectrometer when anal. is complete are achieved by using a signal processor which receives signals from the **NMR** detector or other detectors positioned along the length of the tubing. These signals provide an indication of the position of the sample in the tubing. The signal processor uses this information to adjust the forward and back pressure, thereby achieving the desired positioning of the sample.

NA TAF 6/1/2004